

Respiratory Metabolism

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Introduction: All of the commodities covered in this handbook are alive and carry on processes characteristics of all living things. One of the most important of these is respiratory metabolism. The process of respiration involves combining O_2 in the air with organic molecules in the tissue (usually a sugar) to form various intermediate compounds and eventually CO_2 and water. The energy produced by the series of reactions comprising respiration can be captured as high energy bonds in compounds used by the cell in subsequent reactions, or lost as heat. The energy and organic molecules produced during respiration are used by other metabolic processes to maintain the health of the commodity. Heat produced during respiration is called vital heat and contributes to the refrigeration load that must be considered in designing storage rooms.

There is little the postharvest physiologist can do to alter the internal factors affecting respiration of harvested commodities, since they are largely a function of the commodity itself once harvested. However, a major part of postharvest technology is devoted to reducing respiration and other metabolic reactions associated with quality retention by manipulating the external environment.

In general, the storage life of commodities varies inversely with the rate of respiration. This is because respiration supplies compounds that determine the rate of metabolic processes directly related to quality parameters, eg., firmness, sugar content, aroma, flavor, etc. Commodities and cultivars with higher rates of respiration tend to have shorter storage-life than those with low rates of respiration. Storage life of broccoli, lettuce, peas, spinach, and sweet corn (all of which have high respiration rates) is short in comparison to that of apples, cranberries, limes, onions, and potatoes - all of which have low respiration rates (Table 1).

Table 1. Respiration rates of a range of perishable commodities.

Class	Range at 5 °C ($mg\ CO_2\ kg^{-1}\ h^{-1}$)	Commodities
Very Low	< 5	Nuts, dates
Low	5 to 10	Apple, citrus, grape, kiwifruit, onion, potato
Moderate	10 to 20	Apricot, banana, cherry, peach, nectarine, pear, plum, fig, cabbage, carrot, lettuce, pepper, tomato
High	20 to 40	Strawberry, blackberry, raspberry, cauliflower, lima bean, avocado
Very High	40 to 60	Artichoke, snap bean, Brussels sprouts, cut flowers
Extremely High	> 60	Asparagus, broccoli, mushroom, pea, spinach, sweet corn

Factors Affecting Respiration

Respiration is affected by a wide range of environmental factors that include light, chemical stress (eg., fumigants), radiation stress, water stress, growth regulators, and pathogen attack. The most important postharvest factors are temperature, atmospheric composition, and physical stress.

Temperature: Without a doubt, the most important factor affecting postharvest life is temperature. This is because temperature has a profound affect on the rates of biological reactions, eg., metabolism and respiration. Over the physiological range of most crops, ie., 0 to 30 °C (32 to 86 °F), increased temperatures cause an exponential rise in respiration. The Van't Hoff Rule states that the velocity of a biological reaction increases 2 to 3-fold for every 10 °C (18 °F) rise in temperature.

The temperature quotient for a 10 °C interval is called the Q_{10} . The Q_{10} can be calculated by dividing the reaction rate at a higher temperature by the rate at a 10 °C lower temperature, i.e., $Q_{10} = R_2/R_1$. The temperature quotient is useful because it allows us to calculate the respiration rates at one temperature from a known rate at another temperature. However, the respiration rate does not follow ideal behavior, and the Q_{10} can vary considerably with temperature. At higher temperatures, the Q_{10} is usually smaller than at lower temperatures. Typical figures for Q_{10} are:

Temperature	Q_{10}
0 to 10 °C	2.5 to 4.0
10 to 20 °C	2.0 to 2.5
20 to 30 °C	1.5 to 2.0
30 to 40 °C	1.0 to 1.5

These typical Q_{10} values allow us to construct a table showing the effect of different temperatures on the rates of respiration or deterioration and relative shelf life of a typical perishable commodity (Table 2). This table shows that if a commodity has a mean shelf-life of 13 days at 20 °C it can be stored for as long as 100 days at 0 °C, but will last no more than 4 days at 40 °C.

Table 2. Effect of temperature on rate of deterioration.

Temperature (°C)	Assumed Q_{10}	Relative velocity of deterioration	Relative shelf-life
0	-	1.0	100
10	3.0	3.0	33
20	2.5	7.5	13
30	2.0	15.0	7
40	1.5	22.5	4

Chilling stress - Although respiration is normally reduced at low, but non-freezing temperatures, certain commodities, chiefly those originating in the tropics and subtropics, exhibit abnormal respiration when their temperature falls below 10 to 12 °C (50 to 53.6 °F). Typically the Q_{10} is much higher at these low temperatures for chilling sensitive crops than it would be for chilling tolerant ones. Respiration may increase dramatically at the chilling temperatures or when the commodity is returned to non-chilling temperatures. This enhanced respiration presumably reflects the cells' efforts to detoxify metabolic intermediates that accumulated during chilling, as well as to repair damage to membranes and other sub-cellular structures. Enhanced respiration is only one of many symptoms that signal the onset of chilling injury, an economically important low temperature phenomenon discussed in more detail in a subsequent chapter.

Heat stress - As the temperature rises beyond the physiological range, the rate of increase in respiration falls. It becomes negative as the tissue nears its thermal death point, when metabolism is disorderly and enzyme proteins are denatured. Many tissues can tolerate high temperatures for short periods of time (eg., minutes), and this property is used to advantage in killing surface fungi on some fruits. Continued exposure to high temperatures causes phytotoxic symptoms, and then complete tissue collapse. However, conditioning and heat shocks, i.e., short exposure to potentially injurious temperatures, can modify the tissue's responses to subsequent harmful stresses.

Atmospheric Composition: Adequate O_2 levels are required to maintain aerobic respiration. The exact level of O_2 that reduces respiration while still permitting aerobic respiration varies with commodity. In most crops, O_2 level around 2 to 3% produces a beneficial reduction in the rate of respiration and other metabolic reactions. Levels as low as 1% improve the storage life of some crops, eg., apples, but only when

the storage temperature is optimal. At higher storage temperatures, the demand for ATP may outstrip the supply and promote anaerobic respiration (see section on “Controlled and Modified Atmosphere Storage”). The need for adequate O₂ should be considered in selecting the various postharvest handling procedures, such as waxing and other surface coatings, film wrapping, and packaging. Unintentional modification of the atmosphere, eg., packaging, can result in production of undesirable fermentative products and development of foul odors.

Increasing the CO₂ level around some commodities reduces respiration, delays senescence and retards fungal growth. In low O₂ environments, however, increased CO₂ levels can promote fermentative metabolism. Some commodities tolerate brief (eg., a few days at low temperatures) storage in a pure N₂ atmosphere, or in very high concentrations of CO₂. The biochemical basis of their ability to withstand these atmospheres is unknown.

Physical Stress: Even mild physical stress can perturb respiration, while physical abuse can cause a substantial rise in respiration that is often associated with increased ethylene evolution. The signal produced by physical stress migrates from the site of injury and induces a wide range of physiological changes in adjacent, non-wounded tissue. Some of the more important changes include enhanced respiration, ethylene production, phenolic metabolism and wound healing. Wound-induced respiration is often transitory, lasting a few hours or days. However, in some tissues wounding stimulates developmental changes, eg., promote ripening, that result in a prolonged increase in respiration. Ethylene stimulates respiration and stress-induced ethylene may have many physiological effects on commodities besides stimulating respiration.

Stage of Development: Respiration rates vary among and within commodities. Storage organs such as nuts and tubers have low respiration rates. Tissues with vegetative or floral meristems such as asparagus and broccoli have very high respiration rates. As plant organs mature, their rate of respiration typically declines. This means that commodities harvested during active growth, such as many vegetables and immature fruits, have high respiration rates. Mature fruits, dormant buds and storage organs have relatively low rates.

After harvest, the respiration rate typically declines; slowly in non-climacteric fruits and storage organs, rapidly in vegetative tissues and immature fruits. The rapid decline presumably reflects depletion of respirable substrates that are typically low in such tissues. An important exception to the general decline in respiration following harvest is the rapid and sometimes dramatic rise in respiration during the ripening of climacteric fruit (Fig. 1). This rise, which has been the subject of intense study for many years, normally consists of four distinct phases: 1) pre-climacteric minimum, 2) climacteric rise, 3) climacteric peak, and 4) post-climacteric decline.

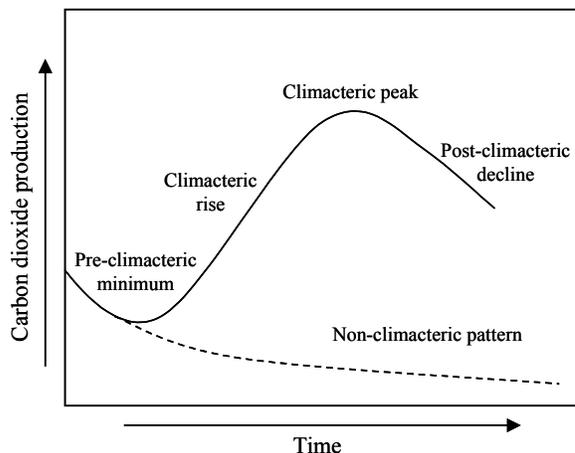


Figure 1. The climacteric pattern of respiration in ripening fruit.

The division of fruits into climacteric and non-climacteric types has been very useful for postharvest physiologists. However, some fruits, for example kiwifruit and cucumber, appear to blur the distinction between the groups. Respiratory rises also occur during stress and other developmental stages, but a true climacteric only occurs coincident with fruit ripening. Following is a general classification of fruits according to their respiratory behavior during ripening:

Climacteric Fruits

Apple
 Apricot
 Avocado
 Banana
 Biriba
 Breadfruit
 Cherimoya
 Feijoa
 Fig
 Guava
 Jackfruit
 Kiwifruit
 Mango
 Muskmelon
 Nectarine

Non-Climacteric Fruits

Papaya
 Passion fruit
 Peach
 Pear
 Persimmon
 Plum
 Sapote
 Soursop
 Tomato
 Watermelon

Blueberry
 Cacao
 Caju
 Cherry
 Cucumber
 Grape
 Grapefruit
 Lemon
 Lime
 Olive
 Orange
 Pepper
 Pineapple
 Strawberry
 Tamarillo

Significance of Respiration

Shelf-life and Respiration Rate: In general, there is an inverse relationship between respiration rates and postharvest-life of fresh commodities. The higher the respiration rate, the more perishable, ie., shorter postharvest-life, the commodity usually is. A summary of respiration and rates is shown in Table 3 of the “General Introduction” section of this Handbook. Respiration plays a major role in the postharvest life of fresh commodities because it reflects the metabolic activity of the tissue that also includes the loss of substrate, the synthesis of new compounds, and the release of heat energy.

Loss of Substrate: Use of various substrates in respiration can result in loss of food reserves in the tissue and loss of taste quality (especially sweetness) and food value to the consumer. For certain commodities that are stored for extended periods of time, such as onions used for dehydrated product, the loss of dry weight due to respiration can be significant. When a hexose sugar (eg., glucose) is the substrate, 180 g of sugars are lost for each 264 g of CO₂ produced by the commodity. The rate of dry weight loss can be estimated as follows: Dry wt loss (g kg⁻¹ h⁻¹) = Respiration (mg CO₂ kg⁻¹ h⁻¹) x 0.068, or % dry wt loss (g 100 g⁻¹ h⁻¹) = Respiration (mg CO₂ kg⁻¹ h⁻¹ x 68 x 10⁻⁶). For example, onions held at 30 °C (86 °F) will respire about 35 mg CO₂ kg⁻¹ h⁻¹. The percent dry wt loss per hour would be 35 x 0.68/10,000 = 0.0024%, while the percent dry wt loss per month would be 0.0024 x 24 x 30 = 1.73%.

Synthesis of New Compounds: Postharvest storage can be used to either prevent any reduction in quality, or to promote changes that increase quality. The quality of most vegetables (eg., cucumbers and lettuce) and non-climacteric fruit (eg., strawberries) is maximal at harvest and storage conditions are optimized to prevent quality loss. In contrast, many flowers (eg., carnations and roses), non-climacteric fruit (eg., lemons and oranges), and climacteric fruit (eg., bananas and tomatoes) are harvested before they reach their best quality and storage conditions are optimized to permit the development of optimum quality. In the first case, the synthesis of new compounds is unnecessary because they lead to reduced quality (eg., enzymes that destroy chlorophyll in lettuce, or promote lignification in asparagus. In the second case, synthesis of pigments and volatiles (eg., lycopene in tomatoes and amyl esters in banana), loss of chlorophyll (eg., chlorophyll degrading enzymes in banana and lemons), and the conversion of starch to sugar (eg., sweetening of apples and bananas) are necessary for development of maximum quality. These synthetic reactions require energy and organic molecules derived from respiration.

Release of Heat Energy: The heat produced by respiration (vital heat), which is about 673 kcal for each mole of sugar (180 g) utilized, can be a major factor in establishing the refrigeration requirements during transport and storage. Vital heat must be considered in selecting proper methods for cooling, package design, method of stacking packages, and refrigerated storage facilities (ie., refrigeration capacity, air circulation, and ventilation). The approximate rates of heat production by various crops at different storage temperatures can be calculated from the summary of respiration rates for many fruits and vegetables given in Table 3 in the “General Introduction” section of this Handbook.

Calculation of heat production from the respiration equation shows that production of 1 mg of CO₂ yields 2.55 cal. In the language of the refrigeration engineer, a respiration rate of 1 mg CO₂ kg⁻¹ h⁻¹ indicates heat production of 61.2 kcal metric tonne⁻¹ day⁻¹ (220 BTU ton⁻¹ day⁻¹). The British thermal unit (BTU) is the heat required to raise 1 lb of water by 1 °F.

Some commodities have high respiration rates and require considerably more refrigeration than more slowly respiring produce to keep them at a specified temperature. For example, asparagus, broccoli, mushrooms and peas respire about 10-times faster than apples, cabbage, lemons, and tomatoes.

Meaning of the Respiratory Quotient (RQ): The composition of a commodity frequently determines which substrates are utilized in respiration and consequently the value of the respiratory quotient (RQ). The RQ is defined as the ratio of CO₂ produced to O₂ consumed; CO₂ and O₂ can be measured in moles or volumes. Depending on the substrate being oxidized, RQ values for fresh commodities range from 0.7 to 1.3 for aerobic respiration. When carbohydrates are being aerobically respired, the RQ is near 1, while it is < 1 for lipids, and > 1 for organic acids. Very high RQ values usually indicate anaerobic respiration in those tissues that produce ethanol. In such tissues, a rapid change in the RQ can be used as an indication of the shift from aerobic to anaerobic respiration.

Measuring the Rate of Respiration: The rate of any reaction can be determined by measuring the rate at which the substrates disappear or the products appear. Apart from the water produced by respiration, which

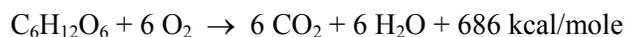
is relatively trivial compared to the very high water content of most harvested commodities, all the substrates and products of respiration have been used to determine the rate of respiration. They are loss of substrate, eg., glucose, loss of O₂, increase in CO₂, and production of heat. The most commonly used method, is to measure production of CO₂ with either a static or dynamic system.

In a static system, the commodity is enclosed in an airtight container and gas samples are taken after sufficient CO₂ has accumulated to be accurately detected by any one of a number of commercially available instruments, eg., gas chromatograph or infrared CO₂ analyzer. If the container is properly sealed, CO₂ should increase linearly with time. Multiplying the change in concentration times the container volume and dividing by weight of the commodity and duration of time between samples gives the production rate.

In the dynamic system a flow of air (or other gas mixture) is passed through the container at a known rate. The system will come into equilibrium (> 99.3%) in about the same time it takes for 5-times the volume to flow through the container. The difference in CO₂ concentration between the inlet and outlet is measured after the system has reached equilibrium by taking gas samples at both points and analyzing them. Multiplying the difference in concentration by the flow rate and dividing by the weight of the commodity is used to calculate the production rate.

Biochemistry of Respiration: Respiration is the oxidative breakdown of complex substrate molecules normally present in plant cells, such as starch, sugars, and organic acids, to simpler molecules such as CO₂ and H₂O. Concomitant with this catabolic reaction is the production of energy and intermediate molecules that are required to sustain the myriad of metabolic reactions essential for the maintenance of cellular organization and membrane integrity of living cells. Since respiration rate is so tightly coupled to the rate of metabolism, measurements of respiration provide an easy, non-destructive means of monitoring the metabolic and physiological state of tissues. For example, events of senescence and ripening are often signaled by abrupt changes in respiration.

Maintaining a supply of high-energy compounds like adenosine triphosphate (ATP), nicotinamide adenine dinucleotide (NADH) and pyrophosphate (PPi) is a primary function of respiration. The overall process of aerobic respiration involves regeneration of ATP from ADP (adenosine diphosphate) and P_i (inorganic phosphate) with release of CO₂ and H₂O. If glucose is used as substrate, the overall equation for respiration can be written as follows:



The components of this reaction have various sources and destinations. The one mole of glucose (180 g) can come from stored simple sugars like glucose and sucrose or complex polysaccharides like starch. Fats and proteins can also provide substrates for respiration, but their derivatives, ie., fatty acids, glycerol and amino acids, enter at later stages in the overall process and as smaller, partially metabolized molecules.

The 192 g of O₂ (6 moles x 32 g mol⁻¹) used to oxidize the 1 mole of glucose diffuses into the tissue from the surrounding atmosphere, while the 6 moles of CO₂ (264 g) diffuses out of the tissue. The 6 moles of H₂O (108 g) that are produced are simply incorporated into the aqueous solution of the cell.

There are three fates for the energy (686 kcal mol⁻¹) released by aerobic respiration. Around 13 kcal is lost due to the increase in entropy (disorder) when the complex glucose molecule is broken down into simpler molecules. Of the remaining 673 kcal that are capable of doing work, around 281 kcal (about 41% of the total energy) is used to produce 38 ATP molecules (38 ATP x 7.4 kcal/ATP). The remaining 392 kcal (57%) is lost as heat. In actuality, most energy is lost as heat since energy is lost to heat every time energy is transferred during a metabolic reaction.

Aerobic respiration involves a series of three complex reactions, each of which is catalyzed by a number of specific enzymes which either: 1) add an energy containing phosphate group to the substrate molecule; 2) rearrange the molecule; or 3) breaks down the molecule to a simpler one. The three interconnected metabolic pathways are glycolysis, tricarboxylic acid (TCA) cycle, and electron transport system.

Glycolysis, ie., the breakdown or lyse of glucose, occurs in the cytoplasm of the cell. It involves the

production of two molecules of pyruvate from each molecule of glucose. Each of the 10 distinct, sequential reactions in glycolysis is catalyzed by one enzyme. Two key enzymes in glycolysis are phosphofructokinase (PFK) and pyruvate kinase (PK). Cells can control their rate of energy production by altering the rate of glycolysis, primarily through controlling PFK and PK activity. One of the products of respiration, ATP, is used in as a negative feed-back inhibitor to control the activity of PFK. Glycolysis produces two molecules of ATP and two molecules of NADH from the breakdown of each molecule of glucose.

Tricarboxylic acid (TCA) cycle, which occurs in the mitochondrial matrix, involves the breakdown of pyruvate into CO₂ in nine sequential, enzymatic reactions. Pyruvate is decarboxylated (loses CO₂) to form acetate that condenses with a co-enzyme to form acetyl CoA. This compound then enters the cycle by condensation with oxaloacetate to form citric acid. Citric acid has three carboxy groups from which the cycle derives its name. Through a series of seven successive rearrangements, oxidations and decarboxylations, citric acid is converted back into oxaloacetate that is then ready to accept another acetyl CoA molecule. Besides producing the many small molecules that are used in the synthetic reactions of the cell, the TCA cycle also produces one molecule of flavin adenine dinucleotide (FADH₂) and four molecules of NADH for each molecule of pyruvate metabolized.

Electron transport system, which occurs on membranes in the mitochondria, involves the production of ATP from the high energy intermediates FADH₂ and NADH. The energy contained in a molecule of NADH or FADH₂ is more than is needed for most cellular processes. In a series of reactions, one NADH molecule produces three ATP molecules, while one FADH molecule produces two ATP molecules. The production of ATP is not only dependent on the energy contained in NADH and FADH₂, but also on the chemical environment (ie., pH and ion concentrations) within the cell and mitochondria.

In the absence of O₂, NADH and FADH₂ accumulate in the reduced form, and as the oxidized forms, (ie., NAD⁺ and FAD, are consumed, the TCA cycle comes to a halt and glycolysis becomes the sole source of ATP production. Regeneration of NAD⁺ is absolutely essential for the survival of the anaerobic cell and takes place during the reductive decarboxylation of pyruvate to ethanol in fermentative metabolism.

Fermentation, or anaerobic respiration, involves the conversion of hexose sugars into alcohol and CO₂ in the absence of O₂. Pyruvate produced through glycolysis via a series of reactions that do not require O₂ can be converted to lactic acid, malic acid, acetyl-CoA, or acetaldehyde. The pathway chosen depends on cellular pH, prior stresses, and the current metabolic needs of the cell. Acidification of the cytoplasm enhances the activity of pyruvic decarboxylase that then shunts pyruvate to form CO₂ and acetaldehyde. The acetaldehyde is converted by the enzyme alcohol dehydrogenase to ethanol with the regeneration of NAD⁺. Two molecules of ATP and 21 kcal of heat energy are produced in anaerobic respiration (alcoholic fermentation) from each molecule of glucose. To maintain the supply of ATP at the aerobic rate, 19 times as many glucose molecules would be needed, and glycolysis would increase 19-fold. However, since only two molecules of CO₂ are produced during glycolysis, instead of six during aerobic respiration, the rate of CO₂ production would not increase by 19-fold but only by 6.3-fold, ie., $19 \div 3$. Concomitantly, there would be substantial accumulation of ethanol and smaller amounts of acetaldehyde. However, glycolysis usually only increases 3- to 6-fold.

The O₂ concentration at which a shift from predominately aerobic to predominately anaerobic respiration occurs varies among tissues and is known as the extinction point, the anaerobic compensation point, and the fermentative threshold. Since O₂ concentration at any point in a fruit or vegetable varies due to rates of gas diffusion and respiration, some parts of the commodity may become anaerobic while others remain aerobic.

References:

- Abeles, F.B., P.W. Morgan and M.E. Saltveit. 1992. Ethylene in Plant Biology, 2nd edition, Acad. Press, NY.
- Biale, JB and RE Young. 1981. Respiration and ripening in fruits - retrospect and prospect. In: J. Friend and M.J.C. Rhodes (eds) Recent Advances in the Biochemistry of Fruits and Vegetables; Acad. Press; NY.,

pp 1-39.

- Kays, S.J. 1991. Postharvest Physiology of Perishable Plant Products. Van Nostrand, 532 pp.
- Lopez-Galvez, G., M.E. Saltveit, and M. Cantwell. 1996. Wound-induced phenylalanine ammonia lyase activity: factors affecting its induction and correlation with the quality of minimally processed lettuce. *Postharv. Biol. Technol.* 9:223-233.
- Ryall A.L. and W.J. Lipton. 1979. Handling, transportation and storage of fruits and vegetables, Vol. 1., Vegetables and melon, 2nd edition; AVI Publ. Co., Westport CT, pp 1-13.
- Ryall A.L. and W.T. Pentzer. 1974. Handling, transportation and storage of fruits and vegetables, Vol. 2., Fruits, AVI Pub., Westport CT, pp. 4-12.
- Saltveit, M.E. 1996. Physical and physiological changes in minimally processed fruits and vegetables. In: *Phytochemistry of Fruit and Vegetables*. F.A. Tomás-Barberán (ed) Oxford Univ. Press, pp. 205-220.
- Tomás-Barberán, F.A., J. Loaiza-Velarde, A. Bonfanti, and M.E. Saltveit. 1997. Early wound- and ethylene-induced changes in phenylpropanoid metabolism in harvested lettuce. *J. Amer. Soc. Hort. Sci.* 122(3): 399-404.
- Wills R.H.H., T.H. Lee, D. Graham, W.B. McGlasson, E.G. Hall. 1981. *Postharvest - An introduction to the physiology and handling of fruit and vegetables*. AVI Pub., Westport CT, 163 pp.